

Fig. 1 In vivo determination of the transepidermal water loss (TEWL) following damage to the skin barrier by SDS treatment

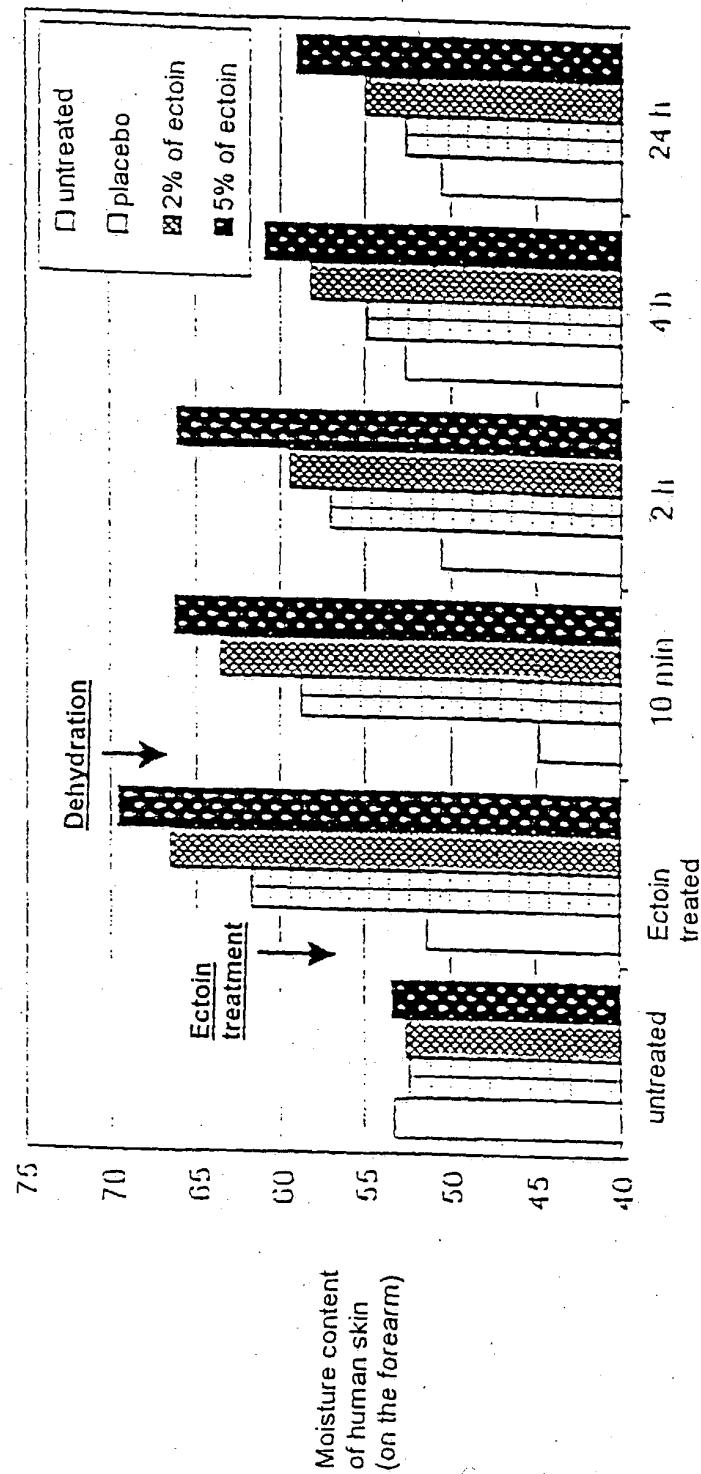


Figure 2: In vivo determination of the skin moisture after ectoin treatment and dehydration by means of silica gel

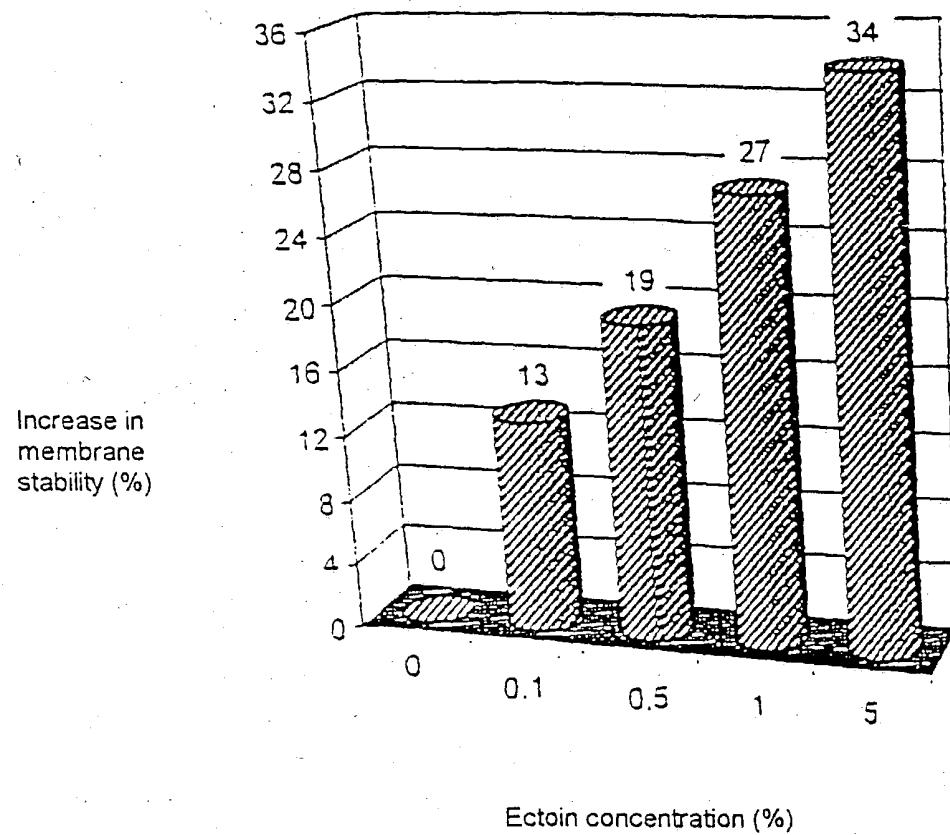


Fig. 3 Determination of the membrane-stabilizing action of human erythrocytes pretreated with ectoin against SDS

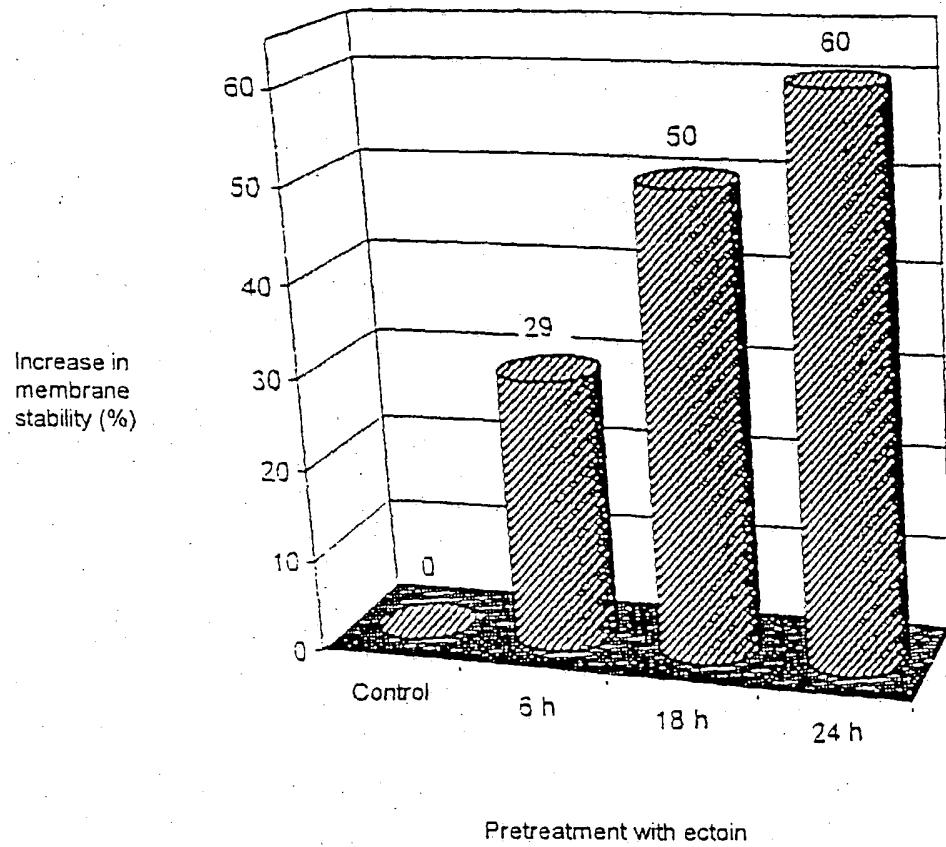


Fig. 4 Determination of the membrane-stabilizing action of human erythrocytes pretreated with ectoin against SDS

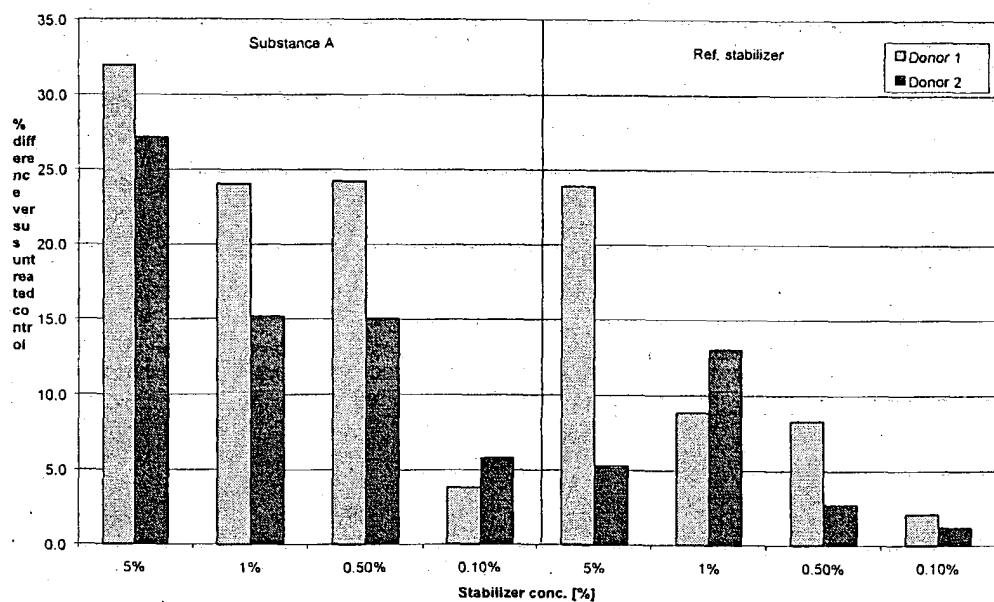
Biosoft D40 H_{50} = 35ppm

Figure 5 : *Determination of the membrane-stabilizing properties of Substance A and reference stabilizer after 1 hour's RBC preincubation. The figure depicts the change in RBC membrane stability (2 donors) versus untreated control. The surfactant H_{50} was used as lytic agent. Data are reported as the mean of 2 assays per protocol and donor.*

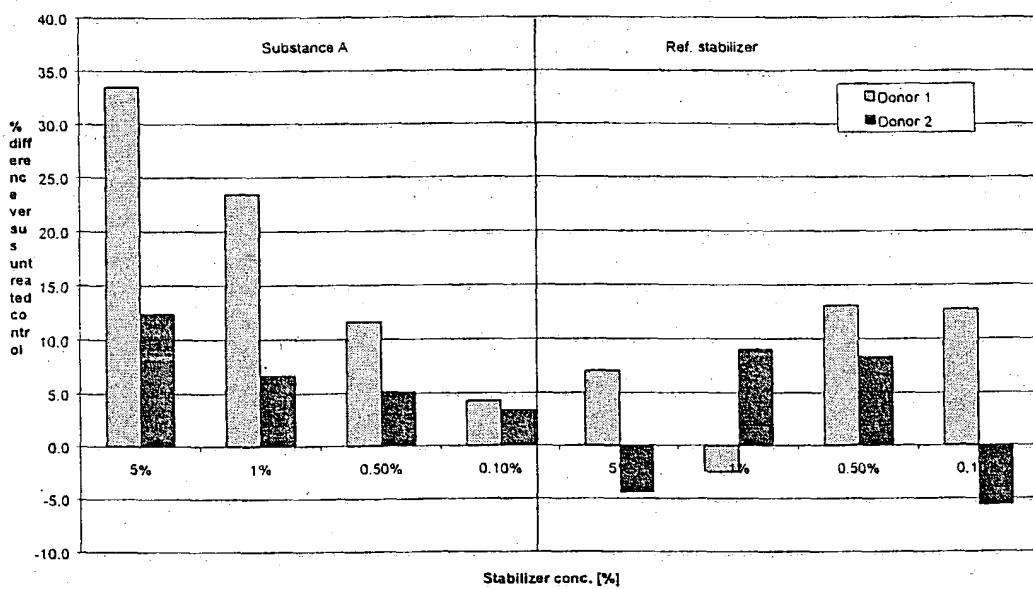
Tego Betain H₅₀ = 65 ppm

Figure 6: Determination of the membrane-stabilizing properties of Substance A and reference stabilizer after 1 hour's RBC preincubation. The figure depicts the change in RBC membrane stability (2 donors) versus untreated control. The surfactant H₅₀ was used as lytic agent. Data are reported as the mean of 2 assays per protocol and donor.

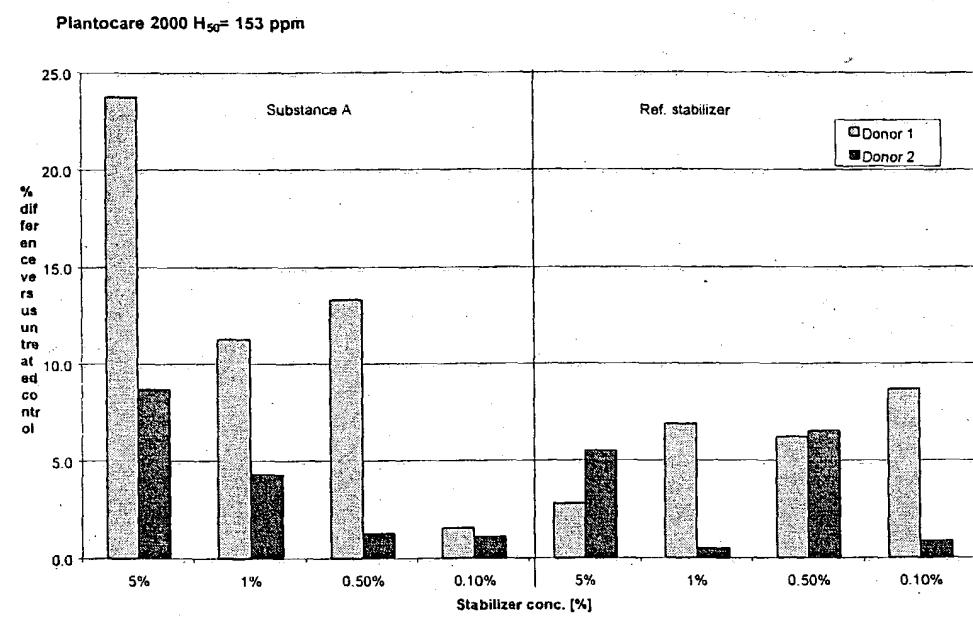


Figure 7: Determination of the membrane-stabilizing properties of Substance A and reference stabilizer after 1 hour's RBC preincubation. The figure depicts the change in RBC membrane stability (2 donors) versus untreated control. The surfactant H_{50} was used as lytic agent. Data are reported as the mean of 2 assays per protocol and donor.

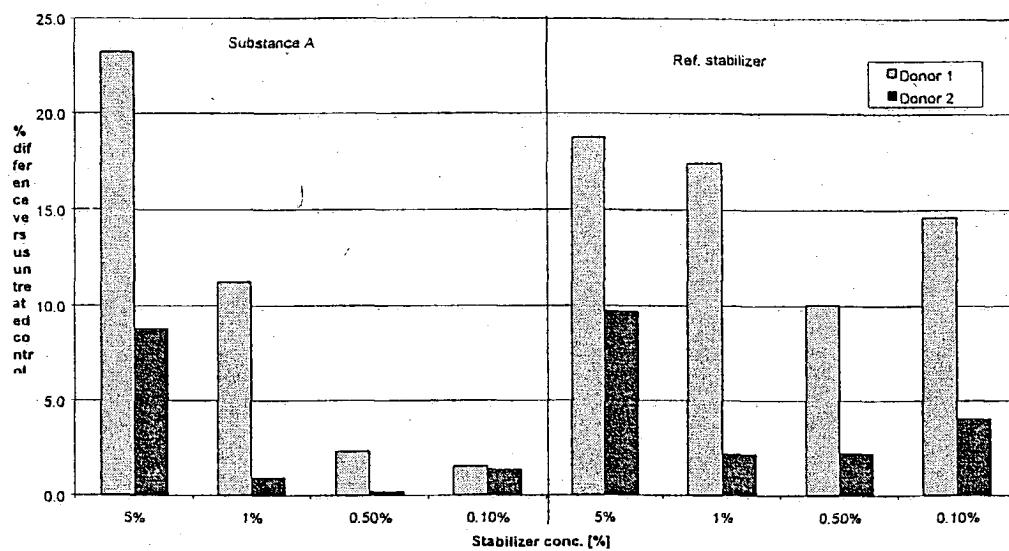
Texapon N SO H_{50} = 44 ppm

Figure 8: Determination of the membrane-stabilizing properties of Substance A and reference stabilizer after 1 hour's RBC preincubation. The figure depicts the change in RBC membrane stability (2 donors) versus untreated control. The surfactant H_{50} was used as lytic agent. Data are reported as the mean of 2 assays per protocol and donor.

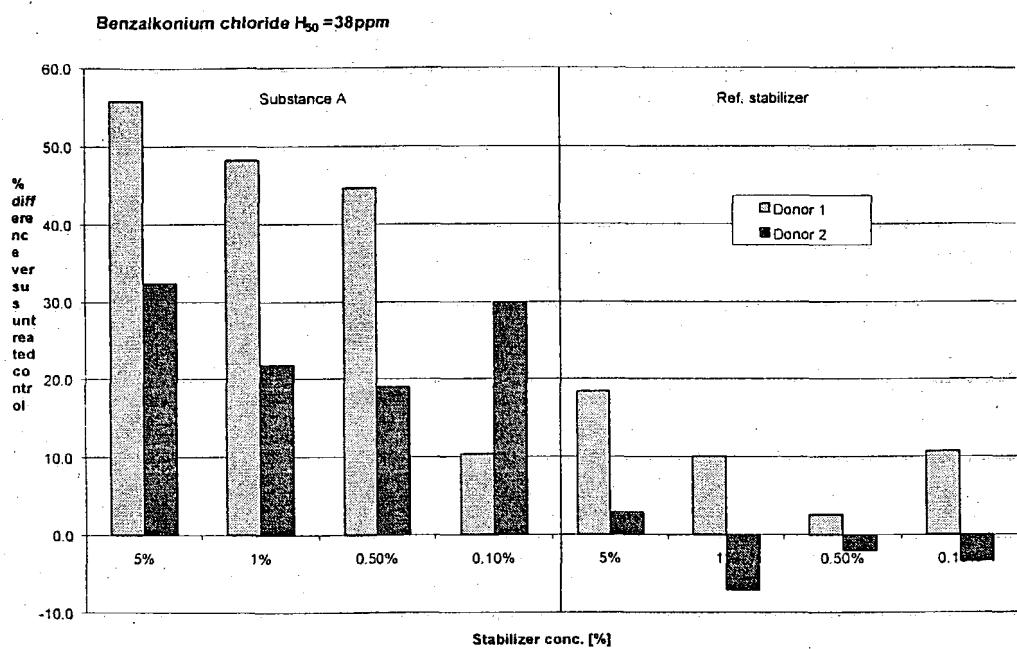


Figure 9 : Determination of the membrane-stabilizing properties of Substance A and reference stabilizer after 1 hour's RBC preincubation. The figure depicts the change in RBC membrane stability (2 donors) versus untreated control. The surfactant H_{50} was used as lytic agent. Data are reported as the mean of 2 assays per protocol and donor.